HEARING

IN THE MATTER OF
PROPOSED AMENDED RULE 111
"CRANIOFACIAL ANOMALY
RECONSTRUCTIVE SURGERY COVERAGE"

HONORABLE DAN HONEY,
HEARING OFFICER
ARKANSAS INSURANCE DEPARTMENT

HEARING PROCEEDINGS
DECEMBER 10, 2021

at 10:00 A.M.

GRIGSBY REPORTING SERVICES
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ALSO PRESENT:

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MS. HOLLY JOHNSON

MS. ASHLI MESARIS

MS. MICHAELA KINLEY

MS. NICOLE HELD

DR. MARK T. JANSEN, M.D.

MS. ZANE A. CHRISMAN

MR. FRANK B. SEWALL

DR. ROBERT TAYLOR

MR. ANDREW TUCKER

MR. GRANT FORTSON

MR. DEAN McNEEL

MS. RITA WILSON, RN

MS. ELLIE WALTON

MR. MILLS BRYANT

MS. SARAH IRONSIDE

MR. MITCH PERSENAIRE

DR. SAGAR MEHTA, M.D.

MS. WENDELYN OSBORNE

MS. BRANDY WEDSTED

MR. JIM BRADER

MS. TERI MECCA

MR. EDDIE DAVIS

MS. CLARA MEZZA

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CAPTION

PROCEEDINGS in the above-styled and numbered cause on the 10th day of December, 2021, before Faith Grigsby, Arkansas Supreme Court Certified Court Reporter #686, at 10:00 a.m., at the Arkansas Insurance Department, 1 Commerce Way, Diamond Mine Hearing Room, Little Rock, Arkansas, pursuant to the agreement hereinafter set forth.

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PROCEEDINGS

DECEMBER 10, 2021

HEARING OFFICER: Good morning. It is

10:00. We are in the Diamond Mine Room 203 at
the Arkansas Department of Commerce, One
Commerce Way in Little Rock, Arkansas. We are
gathered here to consider Arkansas Insurance
Department Proposed Rule 111, Craniofacial
Anomaly Reconstructive Surgery Coverage. I am
Dan Honey. I have been appointed as hearing
officer by Commissioner McClain. The attorney
representing the Department in this matter is
Mr. Booth Rand.

Mr. Rand, would you like to -- do you have
an opening statement?

MR. RAND: Thank you, Mr. Hearing Officer.

I don't, but I would like to admit into the record some exhibits for your consideration at this time.

HEARING OFFICER: Okay. Please proceed.

MR. RAND: Exhibit 1 is the designation of hearing officer form signed by the Commissioner appointing you as hearing officer.

Exhibit 2 is Act 955 of 2021, which is Wendelyn's Law, Craniofacial Law, that

authorizes the rule making that we are engaged in this morning.

Exhibit 3 is the e-mail approval from the Governor's office for issuance of this rule.

Exhibit 4 is a copy of the proposed rule mockup to an earlier Rule 111.

Exhibit Number 5 is a clean version of Amended Rule 111 without the markup.

Exhibit 6 is a copy of the attestation form that has been made part of this rule for purposes of procedures that need to be filed -- or documents that need to be filed by various craniofacial or non-craniofacial providers.

It's part of the rule.

Exhibit Number 7 is a summary that we send to the Arkansas Bureau of Legislative Research.

Exhibit 8 is our Notice of Public Hearing that goes out to the industry by blast, by electronic e-mail, advising the health insurance industry of our hearings. It had a date of November 30, 2021.

Exhibit Number 9 is a copy of Arkansas

Insurance Department Amended Notice of Public

Hearing with a new hearing date of December 10,

2021. We had to do an amendment changing the

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hearing date from November 30 to December 10.

Exhibit Number 10 -- Mr. Hearing Officer, as you know, the Insurance Department has a list of licensees that receive notices of our electronic mail distributions of rules and proposed rules that we do. Exhibit 10 is a copy of the electronic mail distribution to the insurance industry regarding Proposed Rule 111 dated November 1.

Exhibit Number 11 is a copy of another

Notice of Public Hearing. After we changed the hearing date, it was blasted out to the industry.

Exhibit 12 is the first ad we ran at the Arkansas Democrat Gazette advising the public of the date and time of this hearing, and the subject matter of the hearing, which ran October 31st, November 1st, and November 2nd in the classified ads at the back of the newspaper.

Exhibit 13 -- after we changed the hearing date, we ran another ad advising the public of today's hearing date, which ran November 5th, November 6th, and November 7th.

Exhibit 14 is an e-mail and copy of the

Arkansas Insurance Department October 29, 2021, letter to Jessica Whittaker of BLR with the Legislative Council questionnaire, Financial Impact Statement, and Economic Impact Statements.

As the Hearing Officer understands for those, we have to file various numbers of documents with BLR, a lot of it dealing with financial impacts about rules that we're promulgating.

Exhibit 15 is an e-mail to Jessica
Whittaker, again, with updated Notice of Public
Hearing.

The rest of these documents -- Exhibit 16 -- are courtesy copies of filings that we make with various state officials.

Exhibit 16 is a copy of the Arkansas

Insurance Department letter and e-mail to

Secretary of State advising SOS of our proposed rule making.

Exhibit 17 is a copy of the Insurance

Department letter and e-mail to the Secretary

of State with the updated Notice of Public

Hearing.

Exhibit 18 is a courtesy copy that we sent

to the Attorney General's Office advising the Attorney General's Office of our proposed rule making.

And Exhibit 20 is a copy of the Arkansas Insurance Department letter and filing with the Arkansas State Library about the proposed rule.

Exhibit 21 are comments that we received on behalf of the rule during the public comment period.

Exhibit 21, Exhibit Number 1 is a letter in support of the rule by the American Society of Plastic Surgeons.

Exhibit 2 are questions and answers by BLR, Attorney Lacey Johnson, and she engaged with me in the e-mails related to various technical questions related to the rule.

Exhibit Number 3 is a letter in opposition of the rule from Lax Vaughan Fortson Rowe & Threet, by Grant Fortson, who's an attorney who may be speaking here today against the rule.

Exhibit 4 is a letter that we received somewhat advising us of different various issues by Arkansas Blue Cross Blue Shield by Frank Sewall.

They are here today. I don't really want

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to go through all their letters. If they're here to speak, they can bring up those points here when they testify.

HEARING OFFICER: And as usually, the procedure as far as our responses to these comments will be generated after the hearing.

MR. RAND: That's correct. We have to do a post hearing summary for BLR. I will address each of the objections and technical correction requests in that particular document, which will be drafted after this hearing is finished.

HEARING OFFICER: Now, you are not going to be calling any witnesses yourself.

MR. RAND: No.

HEARING OFFICER: Would you like to give us a --

MR. RAND: I'll give a brief update. Absolutely.

HEARING OFFICER: -- brief update and summary of what exactly the rule is meant to -- MR. RAND: Absolutely. Thank you, Mr.

Hearing Officer.

The proposed rule sets up procedures for craniofacial claims consistent with the statute. We are trying to improve craniofacial

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claim adjudications with this rule and with this law by making sure that these craniofacial claims are admitted or denied within two working days for non-urgent cases, and for urgent cases within 24 hours. So the mechanism that we have established is by doing it through this rule.

And as I indicated, one of the things that we wanted to try to clarify was -- and we did create an attestation form for providers for non-craniofacial providers to submit to the ACPA approved teams so that they would have authorization for approval of these services. Those would be submitted to the health benefit plan, and the health benefit plan would only have two working days to actually admit or deny the claim. So we're trying to speed up craniofacial claim adjudications.

The other parts of this rule, we provided definitions of urgent and non-urgent care. The statute did not provide those definitions, and the rule does provide that.

The rule also provides additional coverages for Sclera contact lenses, ocular impressions of eyes, autologous serum eye drops. It

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requires coverage of eye weights and hearing aids, and any incidental tests and procedures.

But the most important part of this rule -if I could identify one of the most important parts of this rule is, we are now going to be requiring health insurers to pay the ACPA approved teams for evaluations and consultations for outside work. So when an ACPA team gets a referral, or a referral is made out to a non-craniofacial provider and the ACPA approved team has to evaluate or consult the actual proposed treatment, the insurance companies now will have to pay the ACPA approved teams for that work. So we believe that's going to financially motivate the ACPA approved teams to do a better job in evaluating and doing consulting and approval process, and help speed up payment of the claims, because they will now be paid.

HEARING OFFICER: I know that we generally don't get into the actual amounts that are paid to providers.

MR. RAND: We're completely --

HEARING OFFICER: Are there going to be quidelines as far as --

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MR. RAND: Yes, sir.

HEARING OFFICER: -- how much they are paid
for these consultations?

MR. RAND: The proposed rule sets out a requirement that the Insurance Department has to issue a bulletin to establish coding fees for these teams. So we will be issuing a bulletin that will establish these fees, and it will be consistent. Every health insurer will have to pay the exact same amount. So Blue Cross won't be paying anything more or less than what United Healthcare would pay. So it's something that we'd have to establish by bulletin.

HEARING OFFICER: It'll be you.

MR. RAND: Yeah.

HEARING OFFICER: Well, if that is all that you have for now, I guess we should -- we have a full house and several folks that would like to testify. I have your list of whether or not you want to testify. I guess it wouldn't hurt to just go down the list and have -- we don't necessarily have the fors first and the against second. So I'll go ahead and start here. It looks like that Michaela Kinley with Ozark

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Prosthodontics. Please come to the --

MS. KINLEY: Good morning everybody.

HEARING OFFICER: Please identify yourself for the record and testify.

MS. KINLEY: My name is Michaela Kinley, like he said. I am a craniofacial patient. I was diagnosed with ectodermal dysplasia, which is a craniofacial anomaly. My brother also has this anomaly. He wasn't able to be here today. And, also, just in advance, if I start to get emotional, I'm sorry. It's just, this has been a really long and hard process that I've walked through.

The thing that's hard is when people see me now and see the -- the main thing is my teeth. They're like, "Well, I don't really understand what you've been through or why this is so hard," but they see the end result, which is what's so great and so easy for me to speak and eat and function and live, but what I had before was not.

So, basically, I was born without the majority of my permanent teeth, and the baby teeth that I have were very small, not very well functioning at all. I couldn't eat a lot.

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My brother couldn't either. They were misshaped. And if you could imagine growing up, that was super hard mentally. And just the ways that we were treated, it was really hard.

And, so, this is my mom, who's Nicole. She went through a lot as a single mom of my brother and I just like figuring out how insurance could help us pay for this because, obviously, we all know it's super expensive. So she spent a lot of time learning about insurance and what Arkansas could provide.

And, thankfully, we are some of the people that were qualified as craniofacial patients because of Act 373, and we were able to -- after a long time -- decades of going through different treatment plans and things failing and not working, and things that were supposed to be the final treatment, but it wasn't working because we just weren't set up with the right provider.

After going through all of this and different, like, legislative things -- my mom knows way more about that than I do. I just remember as a child it was always super stressful. But after going through that long

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process and finally getting set up with the specialty that we needed and getting to see that one day we would be able to have a set of teeth that would help us long term, not something that was just a year fix that might break in a week or something -- which happened all the time, by the way.

But after getting to find these providers that were able to give us a long-term solution and able to utilize this Act for insurance, it was like we finally were able to breathe for the first time in almost -- I mean, I remember this all started when I was 7. I'm 27 now. And I got my final set of teeth when I was 25. So going through that whole process and finally getting to find somewhere that we could trust in the system and had somebody that could provide the best standard of care for us that would last for the rest of my life, I mean, it changed my life forever.

I can tell you, I'm completely different than who I was then. I never really smiled before. I had a lot of confidence issues. And now I'm actually a dental hygienist myself, and I did that because I wanted to provide hope for

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patients like me. And I have felt that I can provide that hope because of what I was able to go through, and that's something that I've always believed in, especially because I have my own daughter and I am super worried that one day she might present with the same symptoms, and that's something that really scares me. And I've always had the hope that this same type of treatment that we have, we could provide for her and utilize insurance.

But now I understand there's some changes that are taking place, and I'm worried that she won't get to utilize that, and that makes me really scared. And just with the families that I've been able to meet along the way have had — felt a lot of hope from seeing my brother and I and what we went through. And so I just hope that you would really consider some of these changes that are being made, because it really affects a lot of people and a lot of families. Sorry. I'm also pregnant again. I'm really emotional, I guess, obviously.

But I also can say that I don't really understand exactly, you know, all the changes that are being made, but I know that there are

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changes. I also used to work at Children's and I know it's a really great place and they want to provide a kind of care. I think some of the proposed plans that are being made, I don't know if the current system, if it's something that can provide the best standard of care for the amount of patients that are needing to utilize insurance for craniofacial anomalies.

So, yeah, I would just hope that you would really consider some of the changes that are being made, you know, thinking long term for my children and for other families that I've met along the way as a hygienist, and just really consider what's being made -- changes. Do you have any questions of me now? Sorry I got emotional. I'm really trying not to.

HEARING OFFICER: Thank you very much for your testimony, Ms. Kinley.

It looks like Nicole Held is next.

MS. HELD: I think she covered it for me.

HEARING OFFICER: Sounds good. I'll call now Mr. Frank Sewall, Arkansas Blue Cross Blue Shield.

MR. SEWALL: My name is Frank Sewall and I'm Senior Counsel of Regulatory Services for

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Arkansas Blue Cross and Blue Shield, and I'm here to speak on behalf of my company in favor of the proposed rule and amended rule.

Arkansas Blue Cross and Blue Shield supports the two principal goals of this rule and the implementing Wendelyn Craniofacial Law.

The first goal is to assure that individuals who are actively diagnosed with a craniofacial anomaly. And the second goal is to assure that patients with a craniofacial anomaly receive appropriate treatment. The proposed rule implements the first goal by requiring, in Section 7, the use of an attestation or authorization form to be signed by the surgical member of an approved craniofacial anomaly cleft palate team when the claim -- and submit it along with the claim to the carrier.

One criticism that I have of the rule is, it only requires so-called non-craniofacial providers. Those are providers who are not part of a craniofacial anomaly team. They're not required under the rule to submit the attestation form. However, the law requires that any craniofacial anomaly, regardless of

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the provider, include attestation or approval by the surgical member of the team.

And we think that's important because this law is unusual in the fact that if the craniofacial anomaly team has diagnosed this person as having a craniofacial anomaly and has approved the treatment plan, the insurance carriers, or the healthcare plans, are very limited in what they can do after that.

Basically, we have to accept the decision and pay for the services. And we're all in favor of doing that because the appropriate people that looked at the patient, according to this rule, and have reviewed the proposed treatment and approved it.

The second goal is implemented by the plan by requiring health carriers to pay the craniofacial anomaly team for its work and making its decision as to whether a craniofacial anomaly exists, and for reviewing the treatment plan, and monitoring the treatment plan as it goes along. We're very much in favor of that. I think this is the only way we're going to get adequate participation by teams, and we're willing to

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work with the Department in helping it propose the annual bulletin.

There are some other changes that I recommended in the rule that are more technical than anything that I could say -- need to say this morning, so I'm going to accept any questions you may have, Mr. Hearing Officer.

HEARING OFFICER: No. I guess specifically, can you address how this -- you think this improves the rule both from the carrier's perspective and from the healthcare consumer's perspective?

MR. SEWALL: I think so. Before the rule and before the recent amendment to the law, the rule -- the law required that a craniofacial anomaly team actually submit its authorization, which would require a review and submission by the team, and there was some -- a lot of confusion as to exactly what that meant. We've had -- we had a number of providers -- or one or two providers that submitted a simple form from a member of the team, or a past member of the team, and that didn't satisfy the terms of the law. So the law was changed to allow a single person on the team to submit the

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authorization form and sign off on it, but it had to be the surgical member; the person that would be doing the actual -- and be a surgical member so that -- because surgery is the most important part of the beginning of the treatment. The other things are -- the other multidisciplinary treatments, the psychology and so forth, are important, but not as important as the surgical.

HEARING OFFICER: Good. Well, it sounds like then that the new law and the new rule -- or amended law and amended rule actually clarify some things and will make the payment of claims run more smoothly from an administrative standpoint. Is that fair to say?

MR. SEWALL: We hope so.

HEARING OFFICER: Do you have any questions, Mr. Rand?

MR. RAND: No.

HEARING OFFICER: No further questions from me. I thank you for your testimony, Mr. Sewall.

MR. SEWALL: You're welcome.

HEARING OFFICER: Mr. Robert Taylor.

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Please identify yourself for the record.

DR. TAYLOR: Dr. Robert Taylor. I'm a craniofacial surgeon in Fayetteville. I had some concerns. You know, one of my largest concerns with the whole rule and the whole law is, it seems to me that it was done in the middle of the night without any input from the providers who actually are on the frontlines taking care of the patients, Number One.

Number Two, I feel like that, when I looked at the law just cursory before I came in, I didn't see anything about acquired craniofacial deformities, yet the rule does mention acquired craniofacial deformities. And from that standpoint as I read the rule, I didn't understand why, as a provider in Northwest Arkansas, I needed to seek approval and confirmation from a provider in Central Arkansas. And Central Arkansas, specifically, Arkansas Children's Hospital, is the only certified cleft lip and palate team in the state of Arkansas.

And the certification for cleft palates and teams has a long past history in that craniofacial surgeons, cleft surgeons are like

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hairdressers. We all think we're the best and we don't think anybody else can do it the same way we can do it. And, so, these teams were designed in the past to jealously guard their territory. So the Dallas cleft lip and palate team, or the North Dallas cleft lip and palate team would seek certification and then seek to block certification of other cleft palate teams. And, so, these teams were grandfathered in to this organization and then these very stringent rules were put in place to prevent further certification of other teams.

And I wouldn't expect the person who wrote the rule to -- and adopted these laws to understand that. But it's important to know that if you can't develop a cleft lip and palate team without having patients, and you can't take care of patients without having a cleft lip and palate team, it makes it almost impossible for other teams to get started.

And then think about the bureaucratic nightmare it's going to require if somebody is in a car wreck in Northwest Arkansas and has panfacial fractures and I have to now seek the -- you know, get the proper documentation, x-

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rays, before I can treat this patient either urgently or semi-urgently, and there's one team -- one person on one team for the entire state of Arkansas, I mean, it's completely untenable. You can't do it.

So I guess that's the biggest reason why I decided to take off work and come down today is to find out why we decided to include acquired craniofacial deformity in a law that was clearly designed for congenital abnormalities. And as a craniofacial surgeon and someone who was on staff at the International Craniofacial Institute in Dallas, who taught other craniofacial surgeons how to do this surgery, I believe in a team approach for developmental disorders and the fact that you need all ancillary services that go along with it, but for acquired deformities, these are things that we have historically taken care of since we -- since medicine has been practiced in Arkansas.

And now, then, unless I'm misreading this rule, it takes the physician on the frontlines diagnostic and treatment protocols and submits them to the whim of someone who maybe was trained with less experience and certainly

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hasn't examined the patient. You can't transport the patient down here to Children's Hospital and have him looked at and then have him transported back to Washington Regional and taken care of.

So I don't see any -- I think the rule is overly broad and I think it's poorly written. I think -- I was looking at Mr. Sewall's -- he thought the reason we needed this law was for accurate diagnosis and appropriate treatment, but I think more likely the law is written for denying of services and limiting access to the patients. I don't see a reason for the acquired deformities, and do applaud the need for teams for taking care of congenital abnormalities.

HEARING OFFICER: Let me ask Mr. Rand a question here.

MR. RAND: I think the problem here -HEARING OFFICER: The acquired -- when
you're talking about there's a 24 hour for an
acute situation such as a car accident --

MR. RAND: The problem that he is pointing his finger to is from the statute itself. The statute of law itself is mandating that these

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authorizations go through an ACPA approved team, and we've only got one in Arkansas. So I'm sympathetic with your problem because it's requiring a funnel to be placed strictly on that ACPA approved team. There's only one in the state, so that team would have to do a fast turnaround on car wreck cases and that sort of thing. So what you're complaining about is from the law itself, not from the rule. So if you want to change this particular process of going through the ACPA approved teams, the best way to do that is through legislation that would repeal the law, or we develop more teams.

So I think your criticism is valid. We have had issues with the ACPA approved teams turning around consultations and evaluations quickly enough for outside providers to do their work. But they've got their own patients at that team itself and they're backloaded, and so one of the things that we wanted to do here is pay them. Make the insurance companies pay them to do consultations and evaluations. So we feel like that might more quickly motivate them to do evaluations for your office. But I can't -- really, I'm sympathetic with what

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you're saying. A lot of it is just institutional.

DR. TAYLOR: Well, let me ask you this, though. When I read the rule -- the law, I didn't see acquired deformities as -- is it part of the law?

MR. RAND: No, it's not. We added that. I will certainly take your comment into -- the Commissioner is going to decide what language is going to be in this proposed rule in response to public comment, so we'll take your public comment and look at that language, for sure.

DR. TAYLOR: And then I think another important thing that people will miss. When you're sitting here -- say I'm an attorney and I think everybody that walks into Children's Hospital and wants to take care of kids is a calling from God. But I'm telling you, when I graduated medical school in Little Rock, there were three programs in the entire country that you could go straight into plastic surgery, and of those three programs, I was lucky to be accepted into two of them.

I decided -- my boss at the time was a

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craniofacial and cleft surgeon. I decided I wanted to do this. It was beautiful surgery. And, so, of the four accepted craniofacial programs, the one in Dallas I was accepted to that, and personally did well enough then that they had asked me to stay on staff and teach other fellows.

I got a call from the chairman of surgery, at the time. He was my -- at the time when I was in medical school. His name was Bob Lawrence. And he said "Bob, I'd like for you to come and apply for the craniofacial position at Children's Hospital." I said, "I'd love to do that, but I'm a cleft surgeon, too." And in 1993, only the ear, nose and throat doctors could do cleft surgery at Children's Hospital. They wouldn't let the plastic surgeons do it. I don't know if that's changed now or not, but I can tell you, in 1993, that was the case. And he said, "Well, you know, otolaryngology is very strong and they're not necessarily under my auspices. I cannot make them let you have those privileges." And I said, "Well, you know, I don't want to come and limit half of my practice just to one thing."

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And so that was my first example, or my first exposure to the fact how jealously this surgery is guarded. And not only that, the International Craniofacial Institute, there's two surgeons there: Ian Munro and Ken Salyer. They hated each other. Vied for patients. I think the biggest problem is, is these deformities, fortunately, are quite rare, and so it takes millions of covered lives in order to generate enough patients to have a practice. And so I can foresee an instance where the grandfathered cleft palate team actively limits the ability for other teams in Jonesboro, El Dorado, Texarkana, Fayetteville, in order to start their own teams, because it might withdraw patients from this central entity.

And, also, I'm curious as to who was in the room when the rules were written, you know.

Was it Arkansas Children's Hospital and Blue

Cross Blue Shield? Because that makes me even

more suspicious of why these rules and this

bottleneck funneling of patients to this one
entity.

MR. RAND: So that's from the law itself, okay? That's not from this rule. If you want

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to change the bottleneck going through ACPA approved teams, you're going to have to change the law. So the way the law is written right now, it's got to go through these teams. So all the rule is doing is amplifying the rule a little bit about what the law talks about. So, again, this process of going through ACPA approved team approval process is something that's in the statute and I can't change.

DR. TAYLOR: Right.

MR. RAND: So if the plastic surgeons and craniofacial providers feel like there's too much of a bottleneck going through these teams to get adequate, timely turnarounds, then that's something you need to visit with, with the legislature.

DR. TAYLOR: But even on a personal level, I wouldn't recognize their authority to even suggest how I took care of a patient. And I think you're going to find that, not just unique to me, it's everybody who's spent the time and the effort into, you know, the expertise in order to do this type of surgery. I think it's poorly written, and I think there were probably nefarious, not just making sure

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that the patients' deformities are covered, but by specifically who covered them and by limiting the amount of access people have to the coverage.

MR. RAND: Again, that's by the law.

DR. TAYLOR: Right. Thank you.

HEARING OFFICER: Thank you, Dr. Taylor. We have Andrew Tucker. Good morning. Please identify yourself for the record.

MR. TUCKER: My name is Andrew Tucker, and I am the CEO of Ozark Prosthodontics. I have worked with many dental practices in my day. In the last eight years I've worked with almost 500 different dental practices, and I had directly worked with at least eight craniofacial team members across the country, predominantly with their orthodontic components. In my time working with those practices, no other practice stands out like Ozark Prosthodontics.

I think it's really important to highlight that Ozark Prosthodontics fits a need in dentistry that doesn't exist in other places.

Ozark Prosthodontics focuses, not on malocclusion, not on cosmetics, but on helping

patients with medical issues resolve those medical issues as they relate to dentistry, doing so under the appropriate licensure in the state of Arkansas, and do so with the appropriate treatment plans and the appropriate standards of care in order to conduct that treatment.

While aesthetic outcomes are certainly part of the equation, there's no crime against that. Plastic surgeons have achieved aesthetic outcomes for reconstruction of their patients for accident victims, for victims of cancer, which is another patient group that we work predominantly with, for years/for decades, and insurance companies have fought over those rules, as well. So we find ourselves kind of in the same group in the same camp.

I highlight all this because we're not the typical dental practice. And I think a lot of times it feels like in the conversations as it relates to this law -- and I promise I'm going somewhere with this. I'm just setting the table. But conversations as it orients around this law feel very accusatory. They suggest that there's some sort of attempt to leverage,

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or some sort of attempt to abuse, or there's an inappropriate standard of care that's being violated. There's attestation requirements that are necessary for treatment. But the reality is, we are what should be, if we could collaborate with the team, an access professional for a craniofacial team. I plan to address that in a moment.

We have a great relationship with the National Federation for Ectodermal Dysplasia. We are on their website in the patient resources. That is one of the most common craniofacial disorders we deal with. And although I'm not a physician nor a dentist, I feel like we are a great place for an ectodermal dysplasia patient to start, like Michaela. We also have a great relationship with the National Organization for Rare Disease, and a great relationship with the Society for Down Syndrome.

Patients that matter tremendously to us, and also patients that are very clearly articulated and carved out in the full spectrum of what we do medically. Patients that are very appropriately treated by our practice, and

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have been historically, and hopefully will continue to be. Although, that's a threat in what we talked about here today.

We are -- like I mentioned, the ACPA, the craniofacial association, we are someone that -- there's not a direct term necessarily we're known for, but we're an access professional.

So, theoretically, a team would leverage us to be able to assist in cases that they have. We have many doctors who have memberships with ACPA, but we are not ourselves in the team. Although, we would love to have that conversation if that was something that was ever on the table.

This isn't to editorialize about the poor legislation that was passed. This isn't to editorialize about how damaging things have been since the passage of the law that fixes to the system have ultimately rendered the system useless. This isn't an editorialization to that. This is really to talk about the core principal of the act and some of the elements in the rule that derive at trying to achieve the primary stated purpose of 955, which is to serve residents in this state who are in need

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of healthcare services to treat craniofacial abnormalities.

Ultimately, when you look at the ACPA and the way teams are structured, there are three required members of an ACPA team. One is a physician, and that physician is not required to be the team lead. One is a dental professional. And the challenge to this is that, in a system that was created -- the ACPA was created and rehabilitated to eliminate a lot of these issues, to create equal footing so people don't practice outside of the scopes of their licensure in order to be able to execute this treatment. In order to pursue that, we have ultimately boiled everything down to the approval of one single individual with an affidavit, and which were ruled out of many teams because -- well, not necessarily many teams, but several teams, because they don't have an M.D. or D.O. or a board certified surgeon.

I've got some important figures to share before I get into the specific issues that we have with the rule. Thirty-one. Thirty-one patients so far in our practice that would have

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been treated under the old format of the rule cannot be treated. These are ectodermal dysplasia patients. These are amelogenesis imperfecta patients. These are patients -- children center patients. These are patients with whom it makes complete sense for treatment to begin in the process of standard of care in the state of Arkansas with appropriate licensure and for us to review.

Things that make complete sense for us to be starting the ball on those cases, we're not permitted to start with those patients, and are not being treated. The patients who are currently in the system while other people wait to be able to be paid for their consults are being paid for on our dime, because the thing we care for the most is seeing that our patients are treated and the treatment is carried out and followed through.

186 teams, at last view -- ACPA teams -- 39 of them do not have an M.D. or a D.O. as a team lead. It's even harder to figure out how many of those are board certified surgeons, but that's a whole different argument in its own right.

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17 of those teams -- 90 percent -- are led by a geneticist. Kaiser Permanente puts geneticists in charge of the teams, because this is considered a genetic team. So this suggestion that surgeons are the most integral part is disagreed with by at least ten percent of the population out there who prioritizes the geneticist, with whom we've typically and traditionally collaborated our very limited cases that we put in for craniofacial purposes, those are the people who collaborated with.

Zero. Zero patients have we been able to collaborate with Arkansas Children's since the tedious and pedantic interpretation of the term "team" that began a year ago. Zero. Zero patients have resulted in collaboration. And zero is the number of times that we have had complaints from patients that treatment plans were inappropriate for patients since the beginning of Ozark Prosthodontics almost 30 years ago.

We fixed a problem that didn't exist. This is a further fix to a problem that doesn't exist to create barriers between the insurance -- create barriers for patients seeking

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insurance reimbursement -- insurance coverage for the treatment that they are contractually and statutorily obligated to. So when it comes to the rule -- we can't fix the legislation today. And when it comes to the rule, I think that a lot of things -- a lot of places, the best that can be done has been done.

But there are really two primary concerns that we have. Mr. Fortson has been kind enough to be able to draft a memorandum that I believe has been entered into the exhibits. But of that -- of the two things I really -- of that whole entire memorandum, I want to highlight two of them in particular. One is the concept of the initiation of a referral for covered services.

I have tremendous concerns about that because the fact that we spent so much time fighting about the definition of what "team" means, and that such a tedious and difficult definition was applied by different parties, I can only imagine that we will all be back in the same room in three months fighting about what the definition of initiated means. I would like to see that we can continue to refer

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patients and, hopefully, that we can see there be a clear of the log jam to be able to actually get them back so that we can pursue treatment for our patients.

Number Two, the attestation of the surgical member. There's a couple of issues with that. Number One, not all oral surgeons are M.D.s or Not all oral surgeons are board certified. That does not mean those people were not competent and capable at their jobs. And the reality is -- a history lesson, in case you're interested -- in 1840, the dentistry department that went to University of Baltimore and said, "Hey, we think we're a medical specialty," and because of the barbaric history of industry as it related to barbers, they said And health insurance companies latched on to that bifurcation and have continued to perpetuate that, despite the direct correlation between healthcare and dental care for the last 150 years. So I say that from the perspective of, I grow frustrated that dental professionals are not viewed in the same respect and the same peerage as physicians.

There are certain cases that are most

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appropriately initiated by physicians. There are other that are most appropriately initiated by dental professionals. For a physician to initiate a dental case first is arguably the unauthorized practice of dentistry, and the statute just put those practitioners in a position to have to do that under this law.

The suggestion that there needs to be notarization on this, in my opinion, is an inference of impropriety. You have people notarize things traditionally in law -- I failed to mention, I'm also a licensed attorney in the state of North Carolina. Traditionally, you have notarization requirements when there is a concern of impropriety and you need someone to affy [sic] as a third party that something wasn't forged.

When we called other craniofacial teams with whom we have a relationship, all of them said, no, we do not have a notary. We don't have a notary. We can go down -- we've got some people who are notaries who do wills and things for people who are kind of, you know, in their dire straits, but we'd have to go down to the ER for that. We'd have to go to, you know,

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intensive care for that. And the next question from many of them was, but why. Why? And it's because, in my opinion, I believe that there is an unfair viewpoint of the role of dentistry in treatment for craniofacial disorders.

Our patients may achieve cosmetic and aesthetic outcomes, but the cases that we initiate are within the scope of our licensure. We feel comfortable with the standard of care to be advising on the things that we are advising on. And why we love collaborating with other physicians, we struggle to find ones that will do it in the boundaries of our own state. This law does nothing but continue to belabor and exacerbate that. Any questions?

HEARING OFFICER: No. Well, I quess, Mr. Rand, I have a question. Everything that you've covered here is included in Mr. Fortson's --

MR. TUCKER: The finer points.

MR. RAND: It's important to note that his physician has also provided us with an alternative language, so Mr. Fortson not only echoed the concerns that this witness raised, but also submitted to the Commissioner

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alternative language for the Commissioner to consider.

HEARING OFFICER: And that will be taken into consideration?

MR. RAND: Yes.

MR. TUCKER: I just want to say, we appreciate the efforts, Booth, that you've had in this rule and hope to find a good one, but we just wanted to raise these concerns.

HEARING OFFICER: Well, thank you, Mr. Tucker.

MR. TUCKER: Thank you

HEARING OFFICER: We have Rita Wilson.

MS. WILSON: Good morning everyone. As he said, my name is Rita Wilson and I have been a patient with Dean, Lori, and Ozark

Prosthodontics since about 2016. I have spent 20 years or more trying to get care that I needed to -- for this congenital abnormality with ectodermal dysplasia and, quite honestly, I'm tired of these fights.

Folks, this is medical. There is no back and forth. You know, I spent all that time -- how many times I walked into a dental office and they would tell me, nothing they could do.

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You talk to the doctor and your doctor tells you, oh no, that's dental. You got to talk to your dental insurance.

I'm here today with someone who has had five family members treated. This starts from the time we're born. The teeth don't develop. The bridges are underdeveloped. I had dentures. What was already underdeveloped just goes away. Somebody had to do something. I went all over looking. It wasn't until, miraculously, I found Ozark and Lori was like, "No, I think we can do something. You need to come see Dean."

Company says we need a form. After all the treatment plan was worked up, that had to be authorized. It had to be sent in to the team. They reviewed it. Now we're talking about sending this form — the way I read the rule, they're talking about, it needs a notary. For what? So we can delay it more? I spent 20 years, folks. I've been doing treatment for five before we got it right. Why are we delaying? Because that's what it seems like this is for is just to put it off. It's one

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more hardship for me to go through; one more thing to fight, you know.

I had this all planned out, but it doesn't really go that way. I get a little upset with it. Honestly, get it right. That's all. This is not something I could have changed. There's nothing we could have done. Acquired or not, when you can't chew food well enough to get it down. When you're worried that you didn't get it chewed up well enough that when you swallow it, it's going to be stuck. Then we go to the ER, right? I've been a nurse for 16 years.

Delaying treatment. Make additional steps. Now we got to have a form. We're going to add more paperwork to it. Anybody remember, we're in a pandemic. Working for the Department of Health, I can tell you, I'm tired, and if you add more paperwork to that, it just makes it that much worse. It's not going to make it better. If you're going to fix this, do it right for the patients. I mean, guys, I travel four hours one way to get treatment.

You want the notary on it. You want it sent it. I'm like Andrew, for what? What's the point of that? And we have standards.

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They're all ethically bound. And I'll take -just like I had to take an oath as a nurse, I'm
going to do my best. I'm going to be a patient
advocate. They all take theirs. But now we're
going to do a notary? To what point? Because
somebody doesn't believe what's being said?

So I'm going to have an extra appointment because we're going to take the time, we got to get the notary, we got to get the provider, we got to get me, get it all scheduled, and I'm going to drive four hours, increasing my cost from that and from the time off work to be able to go up and do it. One more step. One more way to keep me from getting what I need. That much longer.

I'm like Michaela, nobody understands this. You see it. You see it at the end. You didn't see a lot of time coming up with it. You don't see the money that was spent beforehand on what was missing, misshaped, underdeveloped. The \$20,000 that was spent beforehand with braces and the orthodontist and all the other dentists trying to fix it for it to be gone by the age of 30. But then I'm told, it's just a dental problem. No, it's not. No, it's not.

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The fact that I can actually go somewhere now and sit down and eat and I don't have to worry. I can actually eat it. I can actually do it. I know that I'm okay. That's worth something. But without medical insurance covering it, it wouldn't have happened.

Without Dean and Lori helping to help me figure it out and get through it. I've been turned down by more dentists, orthodontists, doctors' offices than I care to admit trying to get it fixed.

So with these rules keep in mind -- I don't know about all the others -- the imperfecta and all the others. I know with ectodermal dysplasia, there's a lot of us out there.

There's a lot of us out there, and every single one of us need the help. I've done everything I could do. I've made calls to senators and representatives to try to get it passed federally so that insurance has to cover.

Arkansas was ahead of the game. I was lucky when I started this, because we were ahead of the game. The state actually had it put in place. There are others in Texas and Oklahoma that don't get that because their

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state doesn't require it. So if we're going to do it, fix it right, please. I think I'm good. Do y'all have any questions?

HEARING OFFICER: None from me.

Do you have any questions?

MR. RAND: Huh-uh.

MS. WILSON: Thank you.

HEARING OFFICER: Thank you. The last name we have signed up here is Mills Bryant.

MR. BRYANT: Hello, my name is Mills
Bryant. I took time out of my day today. I
came from Fayetteville. And in lieu of
studying for my final on Monday -- I've got a
contracts final for my law school class -which, I'm listening to the stuff on the way,
so don't worry. But I came down here because
this is important to me and because I've not
only received competent care from Ozark
Prosthodontics, I received extraordinary care
from Ozark Prosthodontics.

And this rule seems -- the way I read it -I'm not completely versed in this whole deal,
but this rule seems to bar care that I have
received from Ozark Prosthodontics. It seems
to put a railroad block in their way from

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providing me the care that they have. And, to me, that is super important.

I'm obviously a patient of theirs. I've got a genetic disorder. And, literally, me and my wife pray that our children do not have the same disorder. She's already saying, "Mills, I pray they don't have your teeth." And I agree with her. And we say that in jest, but it's true. I mean, the things that we've gone through have been explained today, and they're true. When I was a small child, I couldn't even enjoy ice cream.

You know, I received -- I was a big football player, so people didn't really mess with me, but the one thing they did mess me with was about my teeth, and they would make fun of me and all this kind of stuff saying they looked like corn nuggets and all this kind of stuff. And I took that to heart, because there's nothing I could do. And my parents tried to receive dental care. They were educators and they gave us everything we needed, but they couldn't provide those things for us because dentists would give us exorbitant numbers saying this is what you'll

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have to pay to correctly fix this issue.

And one day we found out about Ozark

Prosthodontics and what they could do for us,
and ever since we found out that day, they have
provided the most extraordinary care that we
could have ever asked for. And now I'm
confident. I have teeth that I can be proud
of. I don't hurt. I don't ache. I can eat
what I want to eat, and things that are
enjoyable.

So this rule seems to block my access to their care, because it's not even clear that I would be able to receive care from them. If the board did not approve that care from Ozark Prosthodontics, I may have to go to someone else which, to me, would be -- I wouldn't do it. I just wouldn't do it. I'd have to find a way around it. And it also -- the rule that I read is too broad giving discretion to the board, because it seems that they could deny care if they didn't think that it was urgent, or if they thought that the non-urgent care that we were asking for was unnecessary.

So, to me, this rule is written not clearly enough. All it says for non-urgent care is

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just care that's not urgent. I mean, if we're going to come up with a definition, we need to do a better job of that. And we need to make sure that patients receive care through insurance. And that's why I'm here today to voice my concerns that I won't be able to receive care, but, in the future, if I do have a child who is unfortunate enough to have this condition, that they won't receive care either. And, you know, we all want to do the best by our kids and, in the future, I want to do the same thing, so it's very important for me to be here today. So thank you very much.

HEARING OFFICER: Thank you, Mr. Bryant.

That is everyone that I have listed to sign up.

Obviously, if there is anyone else in the room

that would like to speak either for or against

the implementation of the rule, please come

forward.

DR. MEHTA: Nice to meet you. My name is Sagar Mehta. I am the director of the craniomaxillofacial team at the Arkansas Children's Hospital. So, for me, you know, I had conversations with Lori McNeel and everyone coming from the get-go. I came here from

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Cincinnati, initially, and I was in Cincinnati for a little bit, and then I was in Salt Lake City before that.

I have done a lot of craniofacial work and it's a passion. Like Dr. Taylor said, it's -- I feel the same way Wendelyn does in some way, where it is kind of like a God's calling. You have these patients that really struggle on a day-to-day basis with who they are, how they feel, and the life that they live, you know. I see that with Michaela. I see that with, you know, many people that have kind of come forth today, and Wendelyn and I have had these conversations before, as well, you know.

It's hard not to bring God or religion or anything into this process because the question is why, right? Why does this happen to certain individuals and not to others, realistically? And my goal and what I do is help those individuals that these congenital and acquired deformities occur for, and to make sure that they are taken care of to the best of our ability.

So, yes, we are the only ACPA approved team in the state. Based on the insurance rules

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that were provided to me before all this stuff went through, I was confined in some ways about how to approve care for Ozark Prosthodontics and things like that. I didn't have the ability in certain instances to make the recommendation, even though I am the surgeon on the team, for them to proceed with care. And I have sent them letters for certain patient populations that they have referred to us.

I am only one man, obviously. And I think that's one of the things that Booth Rand talked about, right, is the manpower that's required in the process of doing this. And because I am a surgical member of the craniomaxillofacial team at Arkansas Children's Hospital, it all kind of falls on me. But I operate on Monday, Tuesday, Wednesday mornings, and Friday afternoon. I am literally coming from a case right now. And, so, I'm only one man.

And I have no argument to the fact that, yes, access to care can be difficult, as Michaela said, and, you know, as Rita has kind of pointed out, as well. It is really difficult, and it's really -- and I will tell you in some ways one of the big reasons why I

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came to Arkansas was because of this law, because I knew that I would just be able to provide care for my patients and not have to battle insurance policies to be able to manage my patients when I know what they need. Does that make sense? And Dr. Taylor is the same way, right, where he knows what they need. And my goal as an ACPA approved team lead in this state is never to impede care.

The rules that had been provided before that, and the conversations that I've had with insurance companies is that the individual that is in question about whether they have a craniofacial anomaly or not is required to see every member of the team. Now, the rule that's set forth before us now is that it has to be the surgical member of the team.

So the realistic thing is that, even though, yes, it does create a funnel on me and/or the partners that I am hiring -- I have two people that we are -- that are interested in coming here, as well, and would be a part of the process, as well. The thing is, is I think that it actually helps to release some of the onus of the dental expectations, or some of

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those expectations that Ozark Prosthodontics is experts at.

I have no qualms with the care that they provide. I really don't. And I think that sometimes, because I kind of sit as the entity, the bubble that kind of gets pushed around between insurance companies and providers and patients, right, in some ways that ultimately, it comes on me. And I'm okay to take that onus and burden. And I try to take that onus and burden with Lori's team and Dean's team, as well as -- we had a meeting about a year ago about this same process.

I'm not here to fight the process. I can only do what insurance companies have requested of me in some ways. So when they say they need to see every member of my team, then I have to bring them into team and I have to have them see every dentist and everyone, and then we have to come up with a team approach and recommendation based off of that. Before that, I wrote you guys a few letters on patients that you guys sent to me, you know, and that wasn't enough, unfortunately. That's not -- that was told to be an inadequate thing. But I actually

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argued that this rule in some ways may help with the expedition of that process, realistically.

And, yes, I won't lie to you in a sense that, when Lori and Dean sent those things to me, again, I'm one man. I have a family of my own. I have two little kids. I have a lot of that stuff. So it is difficult for me to sit down, go through all of the recommendations and all the thought processes, come up with the expertise, read the literature on it, and then kind of come up with a diagnosis, so, for instance, amelogenesis imperfecta.

You know, I was -- I have -- and we have done a lot of literature review and research in some of the things and I -- you know, and I want to create guidelines for each one of these craniomaxillofacial diagnoses. I want to have a checklist that says if they have this, this, this, and this, they have a family history, they have, you know, certain enameloplasty issues, certain measurement, certian this and that, they qualify under amelogenesis imperfecta. That's a checklist. The checklist is checked off. We send them for approval and

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they get approved. Does that make sense?

I want this to be a streamlined process. I'm not here to belabor it or to be the final. I don't want to be the final. It's actually -realistically, being at the end of the final is not really a fun place to be, to be honest. Because, honestly, I want to take care of patients. I feel -- when Rita is talking, and when Michaela are talking, when Wendelyn is talking to me, I see those in my kids, right -the kids that I take care of on a day-to-day basis, and I don't want that for their care. Does that make sense? I don't want them to have to bounce from provider to provider to provider. I want them to find their home. I want their home to then delineate what they need. And that's not --

And the same thing with Dr. Taylor. You know, you expressed with acquired deformities and things of that nature. I don't want to be the final that prevents that patient from getting their facial fractures taken care of in a timely fashion, if that's what's necessary. We all know that there's a time when you can take good care of facial fractures and then

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there's a time where it becomes significantly more difficult. My goal in all of this as the member of the ACPA team that is director is not to impede care in Arkansas -- in the state of Arkansas. It is, in fact, to flourish the care as much as possible and to make it as streamlined a process.

Yes, there are rules and there are issues and there are legislative problems, and there may be -- but the goal of the providers that are in this room, right -- the actual people like Dean and myself and Dr. Taylor, and, you know, I see a whole slew of the Ozark Prosthodontics team here -- should be to rise above that. It shouldn't be about the nuances or the whatever, you know.

And I appreciate Booth's support in some of this. And these are conversations that I actually had with the insurance companies, as well, in terms of, to be able to develop manpower, you need the money. Then we've had some conversation with Dean, as well, you know, where you need money to be able to have people that can help with the processes and expediting things and making sure things happen in a

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timely fashion. And the beauty -- the sum of this rule is that it helps to give us the processes to do so. And I've also brought this forward to the Children's Hospital system, as well.

I want to help the people of Arkansas. And if I -- if people within the state of Arkansas are telling me I've had a congenital condition and I am not being taken care of, you bet your bottom dollar -- I was going to use a more profane word than that -- but you bet your bottom dollar that I will advocate for that patient to the best of my ability.

Wendelyn and I have had conversations hours sometimes and we talk about this, and we talk about the beauty of the process that could be if everything was just -- just happened -- does that makes sense -- and there was no impedance in terms of that process. And I am wholeheartedly about that process, and I will do what I can as the director of the team to make sure those things occur.

I would argue that in some ways, though, yes, this appears like a bottleneck. I don't think it actually is in some ways, because

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rather than this being on the -- I don't disagree with the expertise of Ozark Prosthodontics. I don't disagree with that. But I also -- when they're requested to be met -- be seen by every member of the team, and if a member of the team does not agree with the processes that are in place, I ultimately have to listen to them. I also do believe in the democratic process. I very much believe that everyone has a beautiful mind and they have something to contribute to the world, right. And so for me, personally, whether it's a craniofacial patient or a dental provider or whoever else it may be, I have no arguments with that.

I also believe in accountability. I believe that, you know, I don't want patients that are not "congenital" to have issues in regards to, you know, meth abuse, or whatever it may be, and be covered. And I'm not saying that's what Ozark Prosthodontics is doing, or anyone is doing, realistically. All I'm saying is, is there should be checks and balances in this process, as well.

And I want to help to develop those checks

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and balances with the team members. Like I said, have a checklist. Have things that we, and Ozark Prosthodontics or whatever dental provider may be, agree upon. And when all those things are checked off, then it's not even a question, right. I just have to sign off on the form and make sure that everything occurs the way -- you know, in terms of that.

And that's all I want. I want accountability so that this law doesn't become a beast that can't be controlled in a sense that anyone can get anything done and the onus of that ultimately falls on, you know, whether it's Medicare or Medicaid or any other individual, because that ultimately falls on us in some ways, as well, right.

I want to work with the providers of this team. That's why I came here. That's why I met with you guys. That's why I've had the conversations we've had. That's why I've even sat and spent long hours reviewing your patient charts and doing that. I don't want to be an impedance, and I don't want people like Michaela or Rita, anyone, to feel like they didn't get the care that they deserve. That's

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have. I think I do a good job with the craniomaxillofacial patient population of this state, and I'm proud of the work that we do. I will be absolutely honest with you on that. I do not think that I am the only person in the absolute world that can take care of these

And I don't care. You know, I am a busy

surgeon and I believe in the results that I

just not who I am as a person.

patients and that no one else has the skill or expertise. I know that -- I'm assuming that if

you trained under Ian Munro, who is one of the most forefronts in the craniomaxillofacial

world, then you have a level of expertise.

But you're right, the ACPA has certain burdens associated with it. You have to have a neuropsychologist on board. You have to have all these processes on board. And we have those resources at Arkansas Children's Hospital. And I think it's good. I think some of those patients do need that. I do think if you're talking about the social issues that occur from bullying or from not being able to function and eat for years, or the abuse that some kids go through because they are not

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normal, you know, they need other resources provided to them, and it's really important to provide those resources. And the ACPA has the ability, and we have started working on networking to develop those.

And that's the same thing for dental providers, as well. I don't want this to be a damn. I don't want to block the care and then me pick and choose which ones go through the damn and which ones get stuck. That's not what my goal is in any of this, and I think that's really important for me to stress in this process.

I agree that there are some portions of the law that are frustrating, but, for me, I'm not a legislator and I am not a lawyer. I can assure you, I could have probably been one of those at some point in my life, but that's just not what I chose to be. What I really wanted to be was a patient advocate and a doctor, right. And that's my goal and what I do for my patients is to take care of them and to advocate for them in any way possible, whether that's with Ozark or whether that's with the orthodontists in Jonesboro, or whoever else it

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may be. I will never -- if someone asks me for a craniomaxillofacial patient, how can we get this done sooner, I will do everything in my power to try to make it happen to a fault. You can call my wife if you want to. She tells me I work too much.

So that's where I stand on this. I understand the nuances. I understand that it can be frustrating in some ways. My goal as, again, the only ACPA approved team here within the state is to help support that process. And I am the surgical member of that team, and I honestly think that actually may help in a lot of instances.

Now, everyone is going to have disagreements. That's just the truth of the matter. You expressed that yourself. You know, Ian Munro and Ken Salyer had disagreements, you know. It's not that people don't -- the beauty of a democratic process when everyone contributes to it is, even if there are disagreements, I oftentimes think that that oftentimes allows for the best care of the patient. When one person has absolute control, it doesn't.

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And, yes, it seems like, you know, I'm sitting here and I can delineate one way or the I am not that person. I am never the person that's going to say, "No, this is just what I think. Good bye. See you later." I am very much going to be like, "Well, this is the issue," if there's a question of whether we consider amelogenesis imperfecta -- and I use this term because I think that's one of the diagnosis that we struggled with the most, right. If there's a question whether it's truly a congenital diagnosis or not, which is what has been come back to me about, right, when I talk to insurance companies about whether it is or not, and what true diagnosis amelogenesis imperfecta are not, and what genes they may need to have, or whatever it may be, to get truly approved for that diagnosis.

Again, like I said, all I want to say is,

Ozark Prosthodontics and the ACPA team have

come up with this checklist for what they need

to be diagnosed with amelogenesis imperfecta.

They have checked off all of these

requirements. This is considered a congenital

diagnosis. It needs to be taken care of, and

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that's the end of the matter, right.

Like I don't want to have to fight and I don't want to spend more time than we need to on this. That's just me. I'm sorry. I just — the discrepancy of some of this stuff hurts my soul. I don't like seeing patients cry, like it makes me feel something, too, emotionally. I hate it. The one thing I ask my patients before any case — after any case is, are you happy. And you probably ask the same question in your practice, as well; are you happy with what we did, right.

Because that's the most important thing in this answer, right, is if they're happy with what we did. If they feel better because they can put their best foot forward in that situation, then we've done that for the patient. And that's all I want to do. I want to make sure our patients are happy; they're taken care of appropriately. And that's all. Thank you.

HEARING OFFICER: So you're not necessarily speaking against the rule, but you think that it probably -- both the law and the rule could be improved.

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DR. MEHTA: I don't know the answer. Yes, I guess to some extent there's always improvements that can occur. I think Wendelyn, you know, who had a big part in this rule and this law, did a really good job, right. Like, I am not the only siphon, right. There are other craniomaxillofacial teams that a patient can go to, because the law actually states in there that they can go out of state when they need to, right. So, yes, I am the person that appears in front of you today, but I'm not the siphon to that process. Does that make sense? It could be any ACPA approved team.

And, yes, the politics of that world are old and I don't believe in those politics. I don't believe in the politics that, like, I'm the only one that can take care of craniosynostosis in this state. Do I think I do a really good job with it? Yeah, I'm pretty proud of how I do it.

But I'm not arguing that there aren't other great providers out there that can do the same thing. I'm not arguing that there aren't great providers out there that can do the prosthodontic work part of the population. I

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have no questions from that perspective, and I have no problems with how that works. So, for me, yeah, this is just a law. It doesn't -- again, for me, it means something in the sense that it helps our patient population get the care that they need. But it also -- and it gives me the ability to take care of my patients without a significant impedance in that process. Does that make sense?

DR. TAYLOR: That is exactly the point.

There is no impedance. There is no barrier if the cleft palate team at Children's Hospital decides something needs to be done. But when the Northwest Arkansas cleft palate team comes online, are we going to be reviewing their cases? Are they going to submit their patients and their protocols and then we, therefore, then go over them and give the permission? And how would Dr. Mehta feel about that if, before he could operate on somebody, he'd have to get my permission? I don't think he would be nearly as enthralled with this rule or the law.

MR. RAND: Again, this is --

DR. TAYLOR: This is the law.

MR. RAND: This is the law. And if you

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don't like the law, then go to the legislature and get the law repealed.

DR. TAYLOR: Well, it's not just the law though. The rule incorporates some of that, too, particularly when you require the form be signed.

MR. RAND: I agree. But, again, a lot of the genesis of that is due to the law. So if it gets to a point -- now, I told the Commissioner this -- that our only ACPA approved team is so backlogged and unable to process these claims for evaluation, or they get into business -- or gets into patient load issues, then we need to reevaluate this program. But I'm hoping that -- again, one of the big things this rule does is we're, for the first time, going to pay Dr. Mehta and Children's, or the ACPA approved team to do these evaluations. So I think that will provide some substantiation. There will hopefully be some manpower to do that.

DR. MEHTA: And I also think that your concern about the acquired deformities that occur in patient populations with congenital craniomaxillofacial are -- I never found that

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to be an issue when I was in Cincinnati
(INAUDIBLE) for approval as is with or without
(INAUDIBLE) when I was in Cincinnati. If a
patient had a panfacial fracture, it was
approved. It wasn't like you needed an ACPA
approved team to do so. And I don't imagine
that being an issue, realistically, from my
perspective -- from an insurance coverage
perspective. If that is, then I will fight
with you on that. That's ridiculous. That's
just not how it's supposed to be done.

HEARING OFFICER: That is an acute situation.

DR. MEHTA: That's an acute situation.

DR. TAYLOR: I'm sorry, but I don't see a discrepancy.

MR. RAND: I agree, and it's something we'll look at when we leave out here.

DR. MEHTA: I'm not here to fight with you on that. And I'm not here to -- again, patient access is my primary goal in this process. I never really fought for that otherwise and I -- you know, I know I'm taking a lot of people's time and I apologize. I just feel like I know that there have been a lot of issues in this

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process, but it's more been for me an administrative issue than it has necessarily been a provider issue. And, so, I just want just speak to that before we kind of go through this process.

Yes, everything can change. Everything can be more perfect. But the problem is, is if you get into the nitty-gritties of it, if you start really, really hammering down every single detail, what I'm worried about is, there are some patients that are going to miss out on that process, as well.

HEARING OFFICER: Thank you, Doctor.

DR. MEHTA: Thank you.

MS. McNEEL: Morning, afternoon, whatever it is right now in your time zone. And to the people on Zoom, my name is Lori McNeel. I'm the patient advocate at Ozark Prosthodontics. I happen to be the Lori that people are referencing in the room, just to make sure everybody knows that one. That happens to be my husband, Dr. Dean McNeel. This is year 27 for Dean. I want to be as pertinent and brief as I can for you with Rule 111.

First of all, Children's Hospital can

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absolutely send a claim in for exams, and CTs, and telemedicine, and there's CPT codes for doctor consulting fees. So, Booth, what you're saying with some of this, and Frank, it's not true, and I can't stand here as someone who files medical insurance and look at you guys and say it's fair that you throw the smokescreen up to say they can't get paid for his services. There's 100 CPT codes to do it, and if you can't figure it out, call me. offered that up to you a year ago. These are legitimate CPT codes that the World Health Organization makes up. I didn't invent them. You didn't invent them. There was no reason you couldn't have gotten paid for everything that you reviewed, period. So that is BS, okay.

Number Two, obviously, you've invented a process for this man to be bottlenecked. He's a surgeon trying to help kids three days a week in the operating room. How is he going to do this? This whole fight began because you guys — and I'm seeing one of Arkansas Blue Cross right now — decided that we're a team, but it was your predecessors who invented the process.

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And now where are we? Back to using one person, not the entire team.

I feel as if I'm on Planet Janet, Booth. All team members. Most team members. no. Now a surgical member of the team, and the guy has limited hours in the day. So there's no way in hell this could work. And why you guys fought that, I don't know. And it's unfortunate and embarrassing, and 31 people didn't get help in the state of Arkansas because of your company. You were the ones that fought it, legitimately. And I think every person in the room who has ten minutes of knowledge knows that. So we're reinventing the process that makes it nice for them. Is that what this is about? Because I'm not for that.

I also don't need you to go and get new fee schedules. I can't wait until that disaster happens. Is United going to bring a new contract and, by the way, here's your updated 2022 fee schedules just for the congenital Now, how do I go prove that to them, I'm going to go send them a claim. Booth? you think they've got the ability to decipher that out and re-code all of that and now pay he

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and I just for those craniofacial fees? That's not happening, loud and clear. What are you going to do with an out-of-network doctor? Even if you give them a fee schedule, they haven't hit the out-of-network fees, there is no money taking place. Patients are going to still pay for it.

So why can't we just be honest about what the existing system is and let it work. They've got their fee schedule. If I'm an innetwork doctor -- which I am at this moment; I may not be after this meeting -- but if he's an in-network doctor, we have to live with their fee schedule, no if, ands, or buts. If they allow \$35 an exam, 128 for a CT, then he deserves to get it. We deserve to get it if it's -- you know, period, for any kind of a patient. If it's an allowed service covered in their plan, why are we going to go and get more bureaucracy for that? I think that is a terrible idea, Booth. And I know you were trying to help. I really believe that; AID was really trying to help, Jim, I know that. will not work.

You guys get e-mails from me. You're about

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to get like 30,000. I mean, I don't even want to think about that process, okay.

And you're alluding to the fact that all of these craniofacial teams -- 32 percent of them don't even have an M.D. or a D.O. That's what you put in your form. Your form says only, Frank, no craniofacial surgeon. Only an M.D. or a D.O. board certified can sign off on this form, so 32 percent of teams in the nation can't even play. Now we'll eliminate the ones that don't have a notary on staff. Now we're down to about 26 percent of those that can actually participate. That won't work either.

And, by the way, who's going to pay for the notary? Are we going to give that to the patients to pay for? Are you going to have the physician to pay for it? Are you going to have Arkansas Blue pay for it? Who's paying for the notary? When is the last time you had something notarized? It's somewhere between \$15 and \$35. Well, the patient doesn't have a deductible left. Why would they or Blue Cross pay for that expense? So I'm assuming that would be AID paid for it, because you guys required it.

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MR. RAND: Right.

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MS. McNEEL: Good. Write that down,

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please. MR. RAND: I think we've heard enough

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public comment that --

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MS. McNEEL: About the notary?

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MR. RAND: No, it does not need to be

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notarized. I honestly don't know where that came from. MS. McNEEL: This is the third time we've shut down our practice for this topic. I'm

with Rita. I'm exhausted by it. It makes no

There are good-hearted people, smartminded people in this room to fix problems. If it requires legislation changes, that's happening, okay. Wendelyn, this is your baby. You have to be on point to fix this so patients

opposite way, unfortunately.

Dr. Mehta is doing what he can do, he really is, and he is swamped. I wouldn't expect him to get to read a hello note from any of us Monday through Friday, so you're asking him to do it Friday night through Sunday night, That's not smart. That's not going to right.

can get care. You guys took it just the

work for him.

Last, but not least, with all due respect, if he actually goes forward and does diagnosing planning for us, then I'm going to have to file another lawsuit, because he's operating outside the scope of his license. He can't read enough peer reviewed articles to become an expert.

Dr. McNeel cannot become a plastic surgeon by reading these things. It's a licensure.

That's why we take it so seriously. So let the guys stay in their boundaries where they have these expertise. We are specialists. We don't pretend to be anything else than what -- the way that we file in, right. That's all we want to do.

You have heard these lovely people in the back. They've driven -- Mills, by the way, pray to God he gets a good grade. Everybody pray for that one. But these people are here. You saw them. You heard them, right? These are people who fought for two decades trying to get help.

And we had Arkansas Act 373, and the only thing that went wrong with it was the definition of team, and now you guys have

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flipped the coin over like it didn't happen, and now we're back to one team member can do all this approval. Now, you're going to let a guy that is overwhelmed -- even if he hires 16 more staff members, he doesn't have the licensure or expertise to go do this -- to make these decisions, like we wouldn't for his cases.

MR. RAND: Why didn't you bring this up in legislation? The legislation requires authorization.

MS. McNEEL: We did.

MR. RAND: Okay.

MS. McNEEL: We did. Apparently we got to the game late. But we have full force working on it to ensure it changes. But we have to have a patch between now and then, guys. All you're going to have is now 60 more people showing up mad, 200 more people showing up mad, et cetera.

Let's don't do that. Arkansas is better than this. One thing I was so proud of whenever I go around and talk to the nation about medical insurance and cases like this is how good our state works. We are so lucky.

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We're small and we're lean and we're mean.
We're 49th and we're proud of it. You know
why? Because we fight harder.

So I'm in favor of everyone working together trying to find a simple resolution. So on behalf of what I can say for Ozark Prosthodontics and these people that are in the room with us that flew from North Carolina to be here with you guys to say something, we are here. We're not going away. I think that's clear. We're not going away.

We have -- these people behind me, I'm -Dean and I are going to make sure this happens
one way or the other. I think Dr. Mehta is
committed to his patient base, too. I think
Dr. Taylor is committed to his patient base,
too. Please encourage and get these providers
in these conversations. Please. Please.
Invite some patients in these conversations.
These patients suffer, so please get them. But
Rule 111, bad idea.

MR. RAND: Thank you.

HEARING OFFICER: Thank you. Anyone else?

MS. MILLS: I do have a statement I'd like
to read. Mine is short and sweet. I

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apologize. I was a little bit late getting here. I have to keep it short and sweet and on focus, because this is a topic that is --

HEARING OFFICER: Please identify yourself for the record.

MS. MILLS: My name is Lori Mills. But, again, I have to keep it on focus, because this is a condition I could go on and on about for quite some time.

I have amelogenesis imperfecta, it's a craniofacial anomaly, and my biological sister and her two biological sons also have this condition. My entire life has been continuously getting my mouth worked on.

That's where I could just go on forever. It's been a nightmare. And I've had to deal and struggle with the finances out of pocket.

Having the ability to leverage my health insurance for this treatment is and has been life changing. However, having to jump through hoops year after year is ridiculous.

And I didn't think that, being in my 40s,
I'd have to pause treatment, go to a Children's
Hospital to get an approval, then go back to
finish my treatment. And for the time aspect

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that is difficult enough, it was hard enough for me to get here today. But most importantly, I don't understand how I can pay premiums every month, use providers in my network, but for this part of my treatment I'm told who I can go to for this service. And as I mentioned before, my family has an extensive history with this craniofacial disorder, and finding appropriate providers has been scary. It's been hard.

Had my nephews not taken these first steps, I wouldn't have found the provider that I did, and I have full faith and confidence in them, and I couldn't imagine going anywhere else, and hindering that ability to choose is simply unfair. And I and others with craniofacial anomalies deserve better than this and it's my hope that Rule 111 will be seriously reevaluated after you hear my story.

MR. RAND: Thank you.

HEARING OFFICER: Anyone else?

MS. OSBORNE: Hello everyone. I am
Wendelyn Osborne, the one that you're really
mad at in this room. I'm going to tell you a
little bit about myself, and I'm going to tell

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you, I'm getting -- all of you patients, I'm going to give you some ideas of what you can do, and what we need to do as the state of Arkansas to make this all better.

I started working on this bill -- this law in 2000. 21 years I have been working on this. I started out in D.C. In fact, my very first trip to D.C., when we were going to introduce my bill on the House floor, ended up being -- at 12:00 p.m. ended up being 9-1-1. Osama bin Laden had other plans, so we had to delay and we couldn't do anything until 2002.

While I was in D.C. working on the D.C. bill, I contacted my representative at the town that I was living in at the time, who is also a dentist, and asked if we can write this same bill in the state of Arkansas. I poured my blood, sweat, and tears into getting this law passed for ten whole years. There are lawyers in here. I don't see any legislators in here. You could talk a big game, but if you don't know how to play that game, then it's no good.

I have a craniofacial anomaly myself. In fact, I have the father of craniofacial anomalies as my doctor; Dr. Paul Tessier of

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Paris, France. I had him in the 80s. He did surgeries on me -- Blue Cross Blue Shield -- I've always had an entity of Blue Cross Blue Shield. Always denied it. Always cosmetic. Always cosmetic. Do you know how frustrating that is? It is very frustrating. Do you know how frustrating it is when you're in a lot of pain and you have to walk around and you just want to live normally? You just want to be normal and be accepted, and you're not. I know. I know that. I know some of you had spoken about that this morning. I am well aware of ectodermal dysplasia.

I have craniodiaphyseal dysplasia. I am one -- I was the 16th case in medical history, diagnosed, in the world -- medical history means world -- with craniodiaphyseal dysplasia. I came down with double pneumonia at eight months old. The doctors took an x-ray of my chest and noticed that my bones were just solid white, not shadowy. So they told my parents to go -- I was born on an Air Force base. So they told my parents to go to another Air Force base once I was well, and that's where they diagnosed me as the 16th case in medical

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history with craniodiaphyseal dysplasia. They also told me -- told them that I would not live to be ten and I would never walk. Life expectancy is fourteen.

If you want to get a good whiff of kind of the stuff that I've gone through, go get the movie this weekend, it's called Mask, M-A-S-K, staring Cher. It is the true story of Rocky Dennis. And, by golly, they did a phenomenal job on the makeup on this guy that played Rocky Dennis. And it's so real that I was only -- and I've only been able to watch that one time. And I have -- Cher herself gave me the 25th edition anniversary copy of the DVD. So if you want to get a glimpse of some of the stuff that I've been through.

Just because you have a diagnosis doesn't mean you're going to have every characteristic of that diagnosis, okay. So I've had numerous and numerous and numerous of surgeries. I've had five different sets of braces. I've had -- I can't tell you all the other things that I've had done. But I tell you what, I know pain, I know ignorance, and I know agony on all levels, from lawyers, from

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doctors, from normal human beings just walking around having their own life, okay. We're not all perfect. None of us are perfect. None of us are without pain.

Yes, I'm a little agitated here because some things have been made up that are not true. Don't lie to me or don't lie about me. If you want to get me going, that's what you do. So here's -- I'm going to bring out some things here.

I've had very good doctors. I know craniofacial patients all across the world, and I know them personally, and I help them. know the struggles they go through. I have kids out there. I call -- anybody with a craniofacial anomaly are called my kids. I don't technically have kids. I have kids that are as old as my parents that I call kids. I have kids who have had -- and they're in their -- you know, they have 20-plus surgeries in their single digits years of life. I have kids right now that are in college that have had 49 surgeries, 45 surgeries, okay. It's not fun going under anesthesia. Those of us with craniofacial anomalies tend to have a small

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oral cavity, and it freaks the anesthesiologists out because they're thinking, oh my Gosh, something's going to happen. They freak out. They don't relax.

When you have the conditions -- you know, the craniofacial anomalies that require a lot of surgeries, you become immune to a lot of medications, and your body does not react normal, like they are taught in medical school, and a lot of doctors cannot think outside the box. They have to do textbook-type stuff.

Believe me, I do it to all of them.

Audiologists are hard to find. I wear hearing aids. I have a severe loss and a -- I just went completely blank. I have a severe loss and a profound loss. I have a mixed hearing loss, meaning I have both sensoineural and conductive hearing losses. I even had audiologists tell me that went to Arkansas Medical School, "Oh, well this is your new normal. There's nothing we can do about it." Because of all of the digital hearing aids, everything sounded like Mickey Mouse and Donald Duck.

I'm a vocalist. I sing. I know what real

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sound sounds like. And I looked at some of these audiologists -- well, one particular, and I looked at him and I said, "Hearing aids are a manmade product. Therefore, they can be fixed." I now go to an audiologist who thinks outside the box, got her training outside of the state of Arkansas and, by golly, everything sounds perfectly normal like I've always heard. From the time I was fitted for my very first hearing aid when I was in first grade, six years old.

Okay. So I told you that Paul Tessier was one of my surgeons. I currently have a surgeon that trained under him, and who is world renowned, and I think the world of him. And I know a lot of craniofacial surgeons out on the east coast and I know some on the west. I know some just here in the middle part of the country, like in Tennessee and things like that.

But, you know, we all in this room, we follow rules, okay. Sometimes rules are made up by people, sometimes they're made up by the legislators or laws, et cetera, et cetera. These rules and laws are put into place to

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protect us. You don't run a red light when it turns red, because you could get hit by somebody else. That law is in place to protect us. You may not think of it that way, but that's why. We cannot get through life without rules to help us keep things in order.

Craniofacial anomalies take up two percent of the United States population. In Arkansas we have just over 5,000 living with some form of the craniofacial anomaly. The most common is cleft lip and palate. I worked very hard for 21 years educating federal and state legislators of the ins and outs of living with a craniofacial anomaly. I have not ever benefitted from my law, ever, because I've always had federal insurance. So this being said, I work and I talk with other people -- medical professionals that work in the craniofacial world. They like my law and they like the rule.

Is it perfect? No, it's not perfect, because you have to work with legislators, and they don't understand things all the way through. And the way our legislative session is set up, it's go, go, go, go, go, go, You've

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only got a certain amount of time to get all the stuff done. And I just want y'all to know, our legislators are human beings, too. They have families. They have other jobs that they actually get paid for. And then they're trying to help all the other -- all the people out there, and there's so many people needing things done, they don't have time to get it all done. So try to put yourself in the shoes of the legislators, okay. It's not all about us. They are trying to help. Now, there are some bad ones, just like there are some bad doctors, bad lawyers, bad everything. Okay.

So my law -- I think the world of my kids and there is not one thing I wouldn't do for any of my kids, and I don't even know most of them, okay. My law is put into place to protect the craniofacial patient from well intentioned doctors. That's always been the bottom line. This past legislative session I put forth everything that I know without a doubt in my mind that God told me to put in this amendment and, guess what, there were things that -- even Booth said, "That's never going to happen. That's never going to

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happen." Guess what, it happened. When God wants to do something, he's going to do it and nobody's going to thwart it. No one.

Y'all -- in the law -- in the letter that Mr. Clark wrote, I was just appalled. I'm like, y'all have just basically rewritten this law to say things that it doesn't say. We -- there is nothing in my law that says you have to go to Arkansas Children's Hospital. That is not in my law. You put that in the letter that way. That is not true.

You also state that we need a notary. You don't need a notary. You don't need a notary. You don't have to go pay all that money that Lori says you have to pay, and all the rest of you that think that we have to have a notary, including you.

You know what, we have to work with different people to serve everyone, and that includes our insurance companies. They have to make -- you know, when you're looking at business sides of things, every business has to make money. And I'm going to tell you something right now. Sometimes in life you got to do what you don't want to do. And like Dr.

Sagar -- I'm sorry -- Dr. Mehta said, some things you just don't need to argue about. You just need to figure out how to work together.

The one thing I will not -- I will not -- I will not cave on, there are -- craniofacial teams have rules that they follow. And when I told my doctors -- not just my doctors, but several craniofacial doctors, my law is now, everything begins and ends with the craniofacial surgeons, and they said, thank goodness. All craniofacial teams are busy.

The previous rule that we had -- the former Rule 111 -- because the legislature would not let us go out of the state, and there was only one craniofacial team in Arkansas, and that was Children's Hospital. What about me? Even though I've never benefitted from my law, what about me? I'm an adult and this is my law. But I -- you know, they weren't going to let me benefit from it. Well, we had to write a rule to help those adult patients in the state of Arkansas, and so we did.

But the main thing is, is we put more pressure on the craniofacial team here at Children's, they couldn't do it. They could

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not do it to where they had to monitor medical professionals who had no training in craniofacial anomalies to make sure that the patients were being taken care of and things were done right. Oh, that was -- that just was not a good idea. That was bad.

Now my law specifically says patients can go to any of the ACPA approved team in the United States of America. My law does not say you have to go to Arkansas Children's Hospital, which is what your letter said, Mr. Clark. That is not true. You can -- there -- and the reason I wanted to have that in there is because not every craniofacial team treats every craniofacial anomaly or do they have ample experience in treating that craniofacial anomaly.

I don't know if you remember, or maybe I don't know if I even told y'all this, but I am one of 3,000 in the world with my diagnosis.

It would be hard for you to find a craniofacial team in this country who has one, much less more than that. But the doctor I see, I'm one of six. I'm not going to go to just any craniofacial surgeon whose ACPA approved and

all that, too, if they're not experienced in my condition. You have to have a good rapport with the craniofacial team. Sometimes people just don't get along; they butt heads. You don't need to have that kind of person as your doctor. You have to have respect and you have to be able to trust that doctor with your life or your child's life. There's a right fit for everyone. Currently, at this very moment, there are 200 ACPA approved teams, including Hawaii, if you want to go to Hawaii, that will be covered in-network.

The attestation form is there -- like I told you, my main concern is the safety of my craniofacial kids from well intentioned doctors. That attestation form -- no, it was not my idea but, you know what, I like it. Do you want to know why I like it? Because, no, you don't need a notary, so y'all need to wipe that and scratch that from your little minds, because that's not in -- that is not in the black-and-white writing. Y'all made that up.

In my current law it says the surgical team member -- first of all, if you want to take advantage of my law as a patient or as a

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medical professional, then you need to follow the rules. If you don't, then you don't get to take advantage of it. That's just plain and simple. My law is in place to protect my craniofacial kids from well intentioned doctors.

Believe me, I've had several noncraniofacial doctors well intended that had
made several mistakes on me. I have to live
with them. The doctors can go off and, you
know, live their lives. They don't have to
live with what they've done to somebody, but
the patient does. And pretty much 100 percent
of the time you cannot fully correct what that
doctor did.

Craniofacial doctors on craniofacial teams spend more of their time correcting wonderful procedures done by well intentioned doctors, putting that patient through more surgeries, more pain and all that, and still not be able to fix it. So that's why we've got it in place, everything begins with the surgical team member and ends with the craniofacial surgical team member.

A geneticist does not know how to treat a

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craniofacial anamoly. He might could diagnose it. He might can do all the tests and blood work and all that and tell you what you have, but he doesn't know how -- he or she does not know how to treat it.

Orthodontists can do surgeries, and they
do. Orthodontics is the only area in dentistry
that has a subspecialty in craniofacial
anomalies. Do you want to know why? Because
orthodontists are used in 100 percent of all
craniofacial anomaly cases. Prosthodontists,
endodontists, periodontists and any other
dontist specialist that's in the dental world
are used maybe two percent of all craniofacial
cases. Numbers don't lie, folks.

We're not saying that Ozark Prosthodontics is bad. I don't have anything bad to say about anything that y'all do there. I don't have anything bad to say about that. Just like I don't have anything bad to say about other doctors here in town that I know have done some stuff. But they're not trained. They don't have the proper craniofacial training so, therefore, they need to be monitored by the craniofacial team.

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In fact, the only person who has to sign off on that attestation form is the referring -- the referring, not Dr. -- not the director. Whatever referring surgical team member on that team that recommended you go to see that noncraniofacial person. The thing is, is, they say, okay, well I've done all I can do right now. You need to get such-and-such done. So they have to go see some other subspecialty that they don't have the training in. And, so -- but they could go to anybody they want to in the country, as long as they're Arkansans -they're Arkansas residents, and get it covered, okay. You're not bound to just people in That is stuff that y'all put in your Arkansas. heads. That is not the truth.

But you are going to be -- if you want to take advantage of Wendelyn's Craniofacial Law, by golly, you're going to have to follow the rules. And I don't care if you are a medical professional or if you are a patient. I'm sorry, but I have had enough damage done to me from the well intentioned doctors. I have families -- y'all talk about traveling four hours to go see Dean and Lori. I had families

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that traveled from Vermont down to Dallas. I had families that travel from Ohio down to Dallas, or Colorado down to Dallas. I had families that traveled from New Mexico up to Salt Lake City. I had families who traveled from Florida all the way to Washington State for craniofacial surgeons to get the right craniofacial care that they need for their child or themselves, period.

I'm not denouncing Dr. Sagar at all -- at I'm telling you something. That man loves what he's doing. He has a love and desire for all his patients. And you cannot be a craniofacial surgeon and not love it. Because if you go to a craniofacial surgeon who doesn't like it and is in there for money only -- because let me tell you, you don't make money in craniofacial anomalies, not like you do -- he could go to plastics and make tons of money -- tons more than what he's making. But I'm talking about all the craniofacial surgeons all across the country, they're not in there making all the bookoodles. They could go to the other parts of their profession, whether it's neurosurgery, whether it's orthodontics,

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whether it's otolaryngology, or whether it's plastics, and they can make tons more than what they're making.

So my thing is here. The problem we have in the state of Arkansas -- not in the state of Arkansas, but all over the country, is that obstetrician -- did you know in medical school you might spend like maybe two hours of all your four years of medical school on craniofacial anomalies? So when an obstetrician delivers a baby and it's so deformed -- I think they're all beautiful, by the way -- that is so deformed, they don't know what to do.

The parents -- my parents were first time parents when they had me. I looked like a normal baby. But then my mom -- I was the first grandchild on my mom's side of the family. And at some point my mom asked my grandma, "Well, when is she going to start smiling?" You know, they didn't know until I was ten months old that something was wrong with me, other than the fact that my face didn't move and it was really hard for me to breathe, because my nasal passage was totally

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obstructed.

But -- and I argued -- believe me, I argued with Dr. Paul Tessier in our preop consultation. He said, can you -- "You can't breathe." I said, "Yes, I can." I asked -- I absolutely argued with him. I was in the tenth grade. When I woke up from surgery, I asked my mom, I said -- no, no. I'm sorry. I was in the twelfth grade. He did several surgeries on I said, "There's all this cold stuff in the back of my throat. Why?" She says, "Because you're breathing through your nose." I said, "Oh, I guess Dr. Tessier won that argument." I literally thought I could breathe. Because people with craniofacial anomalies, we've learned to navigate through the things that we can't do. We find other ways to do things.

I've had people -- I'm a vocalist. I've had musical professors that could not get over the fact that my face didn't move and I couldn't do this and I couldn't do that. When we'd go to music lessons and I'm sitting here and they -- we spent the first 15 minutes or 20 minutes of my musical session asking me all

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these questions. Finally, about the fifth time I looked at him and I said, "I don't know how I do anything. This is the way God made me.

Let's get over this. I'm ready to sing."

What we need to do here in the state of Arkansas, we need to educate, get the word out. I don't know what the exact answer is, but we need to get word out to obstetricians. We need to get word out to pediatricians. We need to get word out to everybody so that you and you and you don't have to spend years and years and years looking for treatment when all you had to do was go to a craniofacial team and let them refer you to where you need to go. Let them treat you.

And then if you need to do outside stuff like prosthodontic work, because craniofacial teams do not do prothodontic work -- and obviously -- and what I know for sure is that prosthodontists are needed with ectodermal dysplasia. I have no doubt about that. I don't have a problem with that. But somebody who knows and has many years experience in training craniofacial anomalies and training in those areas needs to be the one to do all the

guiding.

My parents had no clue what a craniofacial anomaly was. It took them six years to find a doctor who'd even look at me, because back in the 60s doctors didn't say "I don't know" like they do now. And then I finally found a doctor -- or they found a doctor that would see me. Guess who she was? Dr. Betty Lowe. She used to be the medical director at Arkansas Children's Hospital. And she got us started, and got us started with the plastics, and the plastics got us started with an orthodontist and an endodontist -- not endodontist, an otolaryngologist, and they created their own craniofacial team before they even existed. Craniofacial teams came in existence in 1989.

So what we need to do is educate, get the word out there so that everybody knows there are craniofacial teams that you need to go to. You cannot under my law -- and it's not safe to do it -- you know, just listen to your friends, or listen to Joe Schmoe, or listen to Google or whatever and take -- pick and choose what you need to do. You don't know all the things about your condition internally. I don't know

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everything about the stuff internally with me.

So you need to have a governing body, which is a craniofacial team -- an ACPA approved craniofacial team. There are 200 in the country. You can go wherever you want to. You don't have to come down here to Arkansas Children's Hospital. You can go wherever you want. And then they do what needs to be done. And then anything that needs to be done outside of their scope and profession, they will tell you where you need -- tell you what you need, and you get to choose who you want to use. But they could also say, well, we know so-and-so and so-and-so. So Lori and Dean could develop a rapport with however many craniofacial teams that they want to and say, "Hey, whenever you need something with prosthodontic work, here we are in Northwest Arkansas." And then when that happens, that's how that could be taken care of.

And then when they go and see Joe Schmoe, well then Joe Schmoe has two days to get back with the referring surgeon on the craniofacial team with what they're thinking about doing. Then they agree on it, you know, whatever.

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They say, okay, that sounds good, blah-blah-blah. And then so that the insurance companies know that these are craniofacial people and so forth and so forth, and keep track so that they don't -- you know, we're trying to protect the craniofacial patient from well intentioned doctors and well intentioned procedures.

It goes back to the referring physician on the craniofacial team -- the referring surgical team member -- to sign that. That is not a notary, and you will not pay 15 or 35 or 1,000 for that notarization. I don't know where in the world y'all came up with that.

And, lastly, I'm going to say this. In 2017, I had surgery. 2015 -- my -- the characteristics of my condition, as you can tell, my seventh nerve -- the seventh nerve is your facial nerve. It controls your facial expressions. I was having -- my bones are still growing. As you're looking at me right now, my bones are still growing. It causes me to have a lot of migraines because there is so much -- my head probably weighs about 40-45 pounds, and I have to carry that on the rest of my body.

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I was having such bad migraines all through 2015 and 2016 that my quality of life was pretty much nothing. I reached out to my surgeon and told him, and he said, go get a CAT scan with contrast. I got the CAT scan with contrast. He consulted the pediatric neurosurgeon on the craniofacial team and they noted that my bone was pressing up so much against all my nerves and everything else in my head, and my brain, which was causing the headaches. So we did a bilateral craniotomy. I call it brain surgery just to be -- make it easier for the layman person to be able to understand, and my surgeon does not like that. Don't tell people that we did brain surgery. But I said it's easier to do that than --

The bilateral craniotomy -- there's several different kinds. But mine, they went in and trimmed away an inch of bone off of each side of my brain to give me some space. But when they went in there, oh my goodness -- these two doctors both have 30 years plus experience in treated craniofacial anomalies of all kinds. Yet again, I'm one of the more rare cases because of my diagnosis. They got in there and

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they were blown away. They had just performed this surgery on a young man the year before that was 20 years old. At the time, I was 50.

Your brain -- if you go to the grocery store today, pick up a cauliflower and feel it and look at the shape of it. That's what your brain is supposed to look like. Mine is smooth as all get-out. The bone was growing everywhere. It had nowhere else to go. Ιt started growing into my brain. So the pediatric neurosurgeon had to intricately take out each of the bone so that it wouldn't cause me to lose all my bodily functions. Because you know, all your bodily functions are controlled in your brain, and there's different areas of your brain. And I'm not going to go into -- I love the brain. That's my favorite part of the body.

So anyway, I am so thankful that I had those doctors doing that surgery on me.

Doctors who know craniofacial anomalies, who have had 30-plus years in working on people with craniofacial anomalies. Each of them.

Not together, each of them. And they had five other people with my particular diagnosis. I

guarantee you, you're not going to find any other surgeon -- craniofacial surgeon out there who will have -- they might have one. I mean, hey, there's -- this is a big world we live in. I've never been outside of the United States other than Toronto, Canada. 3,000 people, that's not very many people when you consider all the people in the world.

I am so thankful that I had them, because they were able to -- even though they were taken back by what they saw, they fell back on all of their training, all their experience with all of their craniofacial experience. And guess what? I am here today. If I had gone to a regular plastic surgeon and a regular pediatric neurosurgeon, I probably would not be here.

When I went back for my two month follow-up postop surgery appointment, I was told that, had I not had my surgery when I did, that my bone growth would have been fatal within five years. And we are monitoring my bone growth annually. And I have so much room in my brain that my doctor told me, he said, "You have all this stuff in your brain to grow." I said,

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"Please don't tell anybody else. They might get a little annoyed, you know, if something else would come up." But, you know, I have a very good memory and stuff like that.

So the things is that we need to do, guys, is, we need to get the word out and start -- and surrounding states, too, to let people know about craniofacial anomalies so when they run into somebody, or somebody runs into them, that they can get them -- say, "You need to go find a craniofacial team." And tell you -- get you the information so that you can go to that team and not have to go and search for this, search for this, and try to treat yourself. You're not trained in craniofacial anomalies. You're not. I'm sorry.

HEARING OFFICER: Could we get back to the topic of the rule or maybe --

MS. OSBORNE: That is the rule. That is the rule.

HEARING OFFICER: (INAUDIBLE)

MS. OSBORNE: And another thing that I'm frustrated with, too. That rule that y'all keep referring to the rule, the rule, the rule, let me tell you something, when you write a

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law, you have to write a rule. It is illegal for us to write stuff in the rule that's not in the law. And you stated that you don't understand why we address the definition for acquired craniofacial anomaly. Yes, that is in the law. Go look at the definition. Go look at the definition of a craniofacial anomaly, and it's in there.

HEARING OFFICER: We're getting close to lunchtime now if you wanted to wrap up --

MS. OSBORNE: I'm done.

HEARING OFFICER: Thank you.

MS. OSBORNE: I'm sorry I got so --

HEARING OFFICER: No, that's fine.

MS. OSBORNE: But the main point is, if you want to take advantage of this law and you -- whether you are a medical provider or you are a patient, then you need to follow the rules.

Otherwise, you don't need to take advantage of it.

HEARING OFFICER: Thank you so much.

MR. TUCKER: I have a point of clarification on the aspects, just in case. The most recent copy of the amendment --

MR. RAND: It is notarized.

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1 MR. TUCKER: I just wanted to make sure --2 MR. RAND: We are not going to require 3 notarization. 4 HEARING OFFICER: Well, we've had much 5 spirited testimony here, and do we have any 6 more? 7 MR. RAND: No. HEARING OFFICER: Anyone else? Mr. Rand --8 9 MR. RAND: The Commissioner will consider 10 everyone's comments. We will get a record of 11 the proceedings from the court reporter and the 12 Commissioner will review your concerns and 13 alternative language and make a decision. 14 HEARING OFFICER: Is the record closed? 15 MR. RAND: The record is closed. 16 HEARING OFFICER: So we will -- and what's 17 our time frame? MR. RAND: Two weeks. 18 19 HEARING OFFICER: Well thank you all very 20 much. We are off the record. 21 (WHEREUPON, the proceedings were concluded 22 in this matter at 12:08 p.m.) 23 24 25

CERTIFICATE

STATE (OF A	ARKANSAS)
)ss
COUNTY	ΟF	PULASKI)

I, Faith Grigsby, CCR, Certified Stenomask
Reporter before whom the foregoing testimony was
taken, do hereby certify that the witness was duly
sworn by me; that the testimony of said witness was
taken by me and was thereafter reduced to typewritten
form under my supervision; that the deposition is a
true and correct record of the testimony given by said
witness; that I am neither counsel for, related to,
nor employed by the parties to the action in which
this deposition was taken, and further, that I am not
a relative or employee of any attorney or counsel
employed by the parties hereto, nor financially
interested in the outcome of this action.

I FURTHER CERTIFY that I have no contract with the parties within this action that affects or has a substantial tendency to affect impartiality, that requires me to relinquish control of an original deposition transcript or copies of the transcript before it is certified and delivered to the custodial attorney, or that requires me to provide any service not made available to all parties to the action.

WITNESS MY HAND AND SEAL this 2nd day of January, 2022.